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**Acetylcysteine for the Treatment of Suspected Remdesivir-Associated Acute Liver Failure in COVID-19: A
Case Series**

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Abstract:

Remdesivir is a direct-acting nucleoside RNA polymerase inhibitor with activity against the novel SARS-CoV-2 virus used in the treatment of COVID-19 pneumonia. Here we present two cases of suspected remdesivir-associated acute liver failure (ALF) in which the liver failure improved following continuous infusion acetylcysteine and withdrawal of remdesivir. Both patients had significant increases in transaminases between day 3 and day 10 of remdesivir therapy accompanied by coagulopathy and encephalopathy. After initiation of continuous infusion acetylcysteine, the transaminases of both patients rapidly improved. Ultimately one patient fully recovered while the other died of suspected septic shock. Due to its novel nature and only recent widespread use, there is very little data on the risk of ALF from remdesivir. Additionally, the data for the use of acetylcysteine to manage non-acetaminophen-induced ALF is limited. It is important to consider the risk of remdesivir-associated ALF when weighing the risk vs. benefits of use, and acetylcysteine may have a role in its management.

Keywords: Adverse drug reactions, Toxicology, COVID-19, Remdesivir, Acute liver failure, Acetylcysteine, Coronavirus, SARS-CoV-2

Background:

The novel severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) was initially identified in Wuhan, China, in December 2019 and has since led to the coronavirus disease 2019 (COVID-19) pandemic experienced worldwide.¹ The first case of COVID-19 pneumonia in the United States was documented in January of 2020, and cases have since exploded to over 5.8 million in the United States with over 180,000 deaths by August 28th, 2020. There are no currently approved treatments for SARS-CoV-2, with recent guidelines recommending supportive care for most patients and the use of remdesivir and dexamethasone for patients with severe COVID-19 pneumonia.^{3,4}

As cases of COVID-19 have increased exponentially, the healthcare infrastructure of the United States has been strained, leading to the desire by many clinicians to use experimental, unapproved therapies in addition to supportive care. One such experimental therapy is remdesivir (Gilead Sciences, Inc, Foster City, CA, U.S.), which is an antiviral with potent activity against SARS-CoV-2.⁵ Remdesivir has been shown to be superior to placebo in shortening time to recovery in adults hospitalized with COVID-19, though there was no difference seen in mortality.⁶ Remdesivir, although not currently approved by the Food and Drug Administration (FDA),

has been made available for clinical use through an Emergency Use Authorization (EUA) program through the FDA. In both pre-clinical and clinical studies, non-serious elevations of both aspartate transaminase (AST) and alanine transaminase (ALT) were noted, and the EUA fact sheet also reports that there have been several cases of severe liver-related laboratory abnormalities.⁷ The EUA program includes monitoring of both AST and ALT as a requirement for use in patients, with a focus on ALT. We report two cases of suspected remdesivir-associated acute liver failure (ALF) in patients with COVID-19 in which the liver failure improved with the addition of continuous infusion acetylcysteine and discontinuation of potentially hepatotoxic medications.

Case #1: A 68-year-old female with a history of hypertension, diabetes mellitus, hyperlipidemia, and coronary artery disease (CAD) presented to an outside hospital with a chief complaint of increasing shortness of breath and tightness across her chest. On admission she was found to have an oxygen saturation in the 70s and a systolic blood pressure >200 mmHg, both treated with bilevel positive airway pressure ventilation (BiPAP) and a nitroglycerin continuous infusion, respectively. The patient also reported chills, fatigue, and body aches for the past few days but with no sick contacts or known COVID-19 exposure. A chest Xray was performed that found bilateral, diffuse interstitial infiltrates consistent with COVID-19 pneumonia, but the COVID-19 reverse transcription polymerase chain reaction (RT-PCR) performed on the day of admission was negative. An initial troponin was collected at a level of 0.2 ng/mL that increased to 1.35 ng/mL over the course of 24 hours, thought to be a type II non-ST-elevation myocardial infarction due to demand ischemia. On day 4 of admission, the patient was taken for cardiac catheterization and was found to have severe multivessel coronary artery disease (CAD), and the patient was transferred to our hospital for coronary artery bypass grafting (CABG). By day 4, the patient had been completely weaned from supplemental oxygen.

On hospital day 6, the patient was taken to the operating room for a planned CABG. Another COVID-19 RT-PCR was sent prior to the OR and was subsequently reported as positive after induction of anesthesia. It was determined that the patient's CAD was critical enough to proceed with the revascularization, and the patient underwent a successful 4-vessel CABG. After the operation, the patient was admitted to the COVID-19 intensive care unit (ICU) where she was extubated to 4 liters nasal cannula without incident on day 7 (post-operative day 1).

The patient's oxygen requirements increased by day 8 to 8 liters nasal cannula and she developed a new cough and worsening infiltrates on chest Xray. A loading dose of remdesivir 200 mg intravenous (IV) was administered the same day due to worsening COVID-19 pneumonia, and a maintenance of remdesivir 100 mg

IV every 24 hours was started on day 9 for a planned total of 5 days of therapy. Prior to remdesivir administration, the patient's liver function enzymes were within normal limits. On day 10, the patient was transferred to the ICU for increasing oxygen demands and an amiodarone infusion was initiated for new-onset, post-operative atrial fibrillation. The amiodarone was initiated with a 150 mg IV loading dose over 10 minutes, and the continuous infusion (450 mg/250 mL) was started at 1 mg/min for 6 hours followed by 0.5 mg/min. By day 10, the patient was requiring humidified, high flow nasal cannula at 60 liters/minute at 100% fraction of inspired oxygen (FiO₂). A complete metabolic panel (CMP) showed an AST >5000 units/L and ALT of 2400 units/L (Figure 1.). Over the course of the next 24 hours, both the AST and ALT were >5000 units/L along with a total bilirubin of 3.1 mg/dL, and International Normalized Ratio (INR) of 2.3, and an ammonia of 161 umol/L. Both remdesivir and amiodarone were discontinued and on the evening of day 10, and a continuous infusion of acetylcysteine was initiated for the suspected remdesivir-induced acute liver failure (ALF) using the 21 hour acetaminophen toxicity protocol of 150 mg/kg over 1 hour, 50 mg/kg over 4 hours, and 100 mg/kg over 16 hours.

By the end of the 21-hour acetylcysteine protocol, the patient's AST and ALT had decreased to 1348 and 1861 units/L, respectively. By day 14, the AST and ALT had decreased to 235 and 966 units/L, respectively, along with a total bilirubin of 2.1 mg/dL and an ammonia of 69 umol/L. Over the course of the rest of her hospital stay, the patient's oxygen requirements improved until she was weaned to room air on day 20 and her LFTs, total bilirubin, and INR returned to within normal limits by day 24 of hospitalization. The patient was then discharged to home with home health.

Case #2: An 80-year-old female with a history of hypertension, diabetes mellitus, and hyperlipidemia presented to the emergency department with low back pain, fatigue, shortness of breath, and difficulty with urination. She endorsed fever and chills but no sick contacts or recent travel. Five days previously, the patient had been admitted to our institution for atypical chest pain and was found to have COVID-19 pneumonia confirmed by RT-PCR, and the patient was subsequently discharged the next day.

Upon this current admission, the patient's chest Xray showed persistent bibasilar and left perihilar infiltrates. She was found to have an oxygen saturation of 82% on room air, initiated on supplemental oxygen of 15 L/minute, admitted to a step-down COVID unit, and started on dexamethasone 6 mg orally every day. By day 3 of admission, she was requiring humidified, high flow nasal cannula at 55 L/minute at 100% FiO₂. Over the next few days, the patient's oxygen requirements remained relatively unchanged. On day 5, she was initiated

on remdesivir, with a 200 mg IV loading dose and 100 mg IV daily for the next 4 days (finishing on hospital day 9). She was also consented for convalescent plasma, which was administered on day 7. While receiving remdesivir, her transaminases remained stable and within normal limits, however, starting on day 5 of remdesivir therapy, her alkaline phosphatase began trending up (but remaining within normal limits during therapy) as well as her total bilirubin.

On day 11, the patient's oxygen requirements continued to escalate, and she was admitted to the ICU where she was placed on BiPAP to maintain her oxygen saturation. She also required norepinephrine and vasopressin continuous infusions due to suspected distributive/septic shock (increasing to a maximum norepinephrine dose of 0.8 mcg/kg/min for around 1 hour on day 13 and then steadily weaned to <0.2 mcg/kg/hr within 24 hours). On day 12 the patient was given tocilizumab 640 mg (7.8 mg/kg) IV once for suspected cytokine storm due to COVID-19. She continued to have worsening respiratory failure, ultimately requiring intubation and mechanical ventilation on day 13. On day 14, the patient had a CMP that showed an AST/ALT of 2178/1510 units/L (Figure 2.), an alkaline phosphatase of 109 units/L, a total bilirubin of 2.1 mg/dL, an INR of 1.7, and an ammonia of 128 umol/L. Due to suspected remdesivir-induced acute liver failure, she was initiated on a continuous infusion of acetylcysteine of 150 mg/kg over 1 hour, 50 mg/kg over 4 hours, and 100 mg/kg over 16 hours. Approximately 12 hours after finishing the 21-hour acetylcysteine protocol, she had an AST/ALT of 546/871 units/L, alkaline phosphatase of 151 units/L, a total bilirubin of 2.3 mg/dL, and an ammonia of 127 umol/L. Her ammonia further decreased to 85 umol/L by 36 hours after finishing the acetylcysteine infusion (a follow up INR was not performed). From day 14 to day 17, the patient's ventilation settings were able to be weaned down from an FiO₂ of 100% to 60%, however, during this time she continued to require vasopressors for distributive shock and went into atrial fibrillation with rapid ventricular rate. On day 17, the patient became more hypotensive requiring reinitiation of a vasopressin infusion, and her arterial blood gas showed a severe metabolic acidosis. Her ventilatory settings escalated back up to requiring 100% FiO₂, and she was found to be hypoglycemic. Despite our efforts, the patient went into cardiac arrest on the evening of day 17 and expired.

Discussion:

It has been reported that infection with SARS-CoV-2 is associated with elevations in transaminases, similar to what has been seen in the original SARS-CoV and Middle East respiratory syndrome coronavirus (MERS-CoV) infections.^{8,9} In most reports, however, transaminases have only been moderately elevated (5-10 times the

upper limit of normal), suggesting liver injury instead of ALF. While it appears that SARS-CoV-2 has some properties that can affect the liver and hepatic function, it is also important to examine if the therapies currently being used for the treatment of COVID-19 may also cause liver dysfunction either independently or in conjunction with the natural course of the virus.

Remdesivir is a novel direct-acting nucleoside RNA polymerase inhibitor with activity against the novel SARS-CoV-2 virus used in the treatment of COVID-19 pneumonia. In Phase 1 clinical trials in healthy volunteers, grade 1 and 2 transaminase elevations were observed which resolved after discontinuation.⁷ In a randomized, double-blind, placebo-controlled trial of 1063 hospitalized patients with COVID-19, there was no difference in the incidence of grade ≥ 3 non-serious elevations of AST, ALT, or both between remdesivir and placebo (4.1% versus 5.9%).⁶ Finally, in a randomized open-label study of remdesivir 5 days vs. 10 days for the treatment of hospitalized patients with COVID-19, the incidence of grade 3 or 4 elevations of AST or ALT ranged between 2-6%.¹⁰ The EUA fact sheet provided by the manufacturer states that in the original compassionate use program of remdesivir in patients with severe or critical illness with COVID-19, liver function test abnormalities were reported in 12% of patients with a time to onset from first dose ranging from 1-16 days. This included cases of serious liver-related laboratory abnormalities.⁷

Outside of experiences discussed in the EUA fact sheet, very little has been published regarding the risk of ALF from remdesivir. One published case report by Leegwater discusses a case of acute liver failure 5 days after initiation of remdesivir in a patient with severe COVID-19 pneumonia.¹¹ In that case, a dramatic increase in the AST and ALT to 1305 units/L and 1461 units/L (respectively) were noted on day 5 of remdesivir therapy. The authors also noted that amiodarone was initiated on day 2 of remdesivir therapy and postulated that amiodarone-induced inhibition of P-glycoprotein efflux transporters (of which remdesivir is a known substrate in vitro) reduces efflux of remdesivir from hepatocytes, increasing remdesivir to toxic concentrations. This is only theoretical, as remdesivir is listed as only a minor substrate of P-glycoprotein efflux transporters and has a relatively short half-life before being converted to active metabolites.¹²

In the two cases we present, the time course of COVID-19 infection, remdesivir administration, and liver function abnormalities appears to point to remdesivir as a possible contributing cause of ALF. When utilizing the Naranjo algorithm to determine the possibility of a drug-induced effect, both cases scored as a “probable” adverse drug reaction with a score of 6 each.¹³ In the first case presented, ALF was noted on day 3 of remdesivir therapy. Though amiodarone can rarely cause ALF, it is unlikely to be the cause in this case as the

ALT was already beginning to trend upward on the day amiodarone was initiated. Also, the ALT began to resolve very rapidly with discontinuation of remdesivir and amiodarone and initiation of acetylcysteine; as amiodarone has a half-life of 9-142 days (depending on formulation and duration of therapy), if amiodarone was the cause of the ALF, it is not expected that discontinuation would lead to such rapid resolution.¹⁴ In the second case, the sudden increase in transaminases was 5 days after completing remdesivir but still within the time range described in the EUA fact sheet. The patient did experience severe distributive shock the day before evidence of overt liver failure, requiring high dose vasopressors for a short time. Hepatic ischemia from distributive shock could have been a contributing factor in development of ALF, but increasing markers of liver injury (increasing alkaline phosphatase several days before vasopressor initiation) prior to the patient experiencing shock along with the timing of the recent remdesivir exposure indicates that there is a possibility of remdesivir also contributing to ALF. The patient also received tocilizumab two days prior to experiencing ALF. Tocilizumab has been associated with mild transaminitis (AST/ALT 1-5 times the upper limit of normal) in acute use that tends to resolve quickly, and there have been cases of severe liver injury linked to tocilizumab use.¹⁵ In these cases, however, severe liver injury was usually associated with several months of tocilizumab therapy, where our patient only received a single dose prior to ALF.

Drug-induced liver injury (DILI) is a serious adverse effect of medications that can range in presentation from mild transaminitis to fulminant liver failure and represents one of the most common causes of acute liver failure.¹⁶ Acetaminophen is the leading cause of ALF in the United States and Europe, though there are many other medications associated with DILI and ALF. Acetylcysteine is the antidote for acetaminophen toxicity, and though acetylcysteine is only indicated for treatment of acetaminophen-induced liver failure, some guidelines recommend the use of acetylcysteine in non-acetaminophen-induced ALF (NAI-ALF) while others recommend its use only in the context of clinical trials.¹⁶⁻¹⁸

Though there are different etiologies of ALF, the loss of hepatocellular function commonly leads to multi-system organ failure if not addressed. While treatment of DILI revolves around stopping the offending agent, other supportive measures are often needed, including administration of an antidote if available, achieving hemodynamic stability, reversing coagulopathies, treating encephalopathies, and consideration of liver transplantation.¹⁹ While acetylcysteine acts as a glutathione precursor in acetaminophen toxicity, its mechanism of action in NAI-ALF is not as clear, though it is hypothesized that acetylcysteine's antioxidant and vasodilatory properties may provide some benefit.¹⁹

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Acetylcysteine use in NAI-ALF is controversial due to the lack of data conclusively showing benefits. Initial trials assessing the use of acetylcysteine in NAI-ALF in adults found conflicting results. A prospective, randomized, double-blind trial by Lee et al. found higher transplant-free survival only in patients with coma grades I-II (representing earlier stage liver failure) treated with intravenous (IV) acetylcysteine compared with placebo.²⁰ This contrasts with a study by Mumtaz et al. who conducted a prospective trial assessing an oral acetylcysteine protocol compared with a historical control group. While the primary cause of ALF in this trial was acute hepatitis, the authors found that not using acetylcysteine was an independent predictor of mortality while also reporting no significant adverse effects from its use.²¹ Subsequent trials in adults with NAI-ALF have continued to support the hypothesis of benefit from acetylcysteine, though they have been retrospective or small. Darweesh et al. conducted a prospective observational trial with historical control of adults admitted with NAI-ALF. They found that patients given IV acetylcysteine on admission had a significant improvement in transplant-free survival (96.4% vs. 23.3% in the placebo group) along with reduction in hospital length of stay, bleeding, and encephalopathy in the acetylcysteine group.²² These results show a much more significant benefit than previously seen, though this could be due to the small, observational nature of the trial. Finally, Nabi et al. conducted a prospective, randomized trial comparing IV acetylcysteine to placebo for treatment of NAI-ALF. The authors found that the acetylcysteine group had significantly lower mortality and hospital length of stay compared to the placebo group and that not using acetylcysteine was an independent risk factor for death.²³ It is important to note that suspected DILI only accounted for anywhere from 6-40% of NAI-ALF cases in the different studies, indicating the difficulty in applying the limited literature to a case of suspected ALF from a novel, unapproved medication used in an emergency setting.

To date, it remains unclear what the true risk and prevalence of ALF from remdesivir use is in the setting of COVID-19. Remdesivir has been studied in humans previously for the treatment of the Ebola virus, but it has only been widely available for clinical use during the SARS-CoV-2 pandemic. While remdesivir has been shown to be relatively safe in the context of limited use, there is a possibility that widespread use will reveal new safety concerns that must be accounted for when weighing the risks vs. benefits of use. The incidence of transaminitis seen in previous clinical trials of remdesivir suggests there is a possibility of unexpected ALF. While there is very limited evidence supporting the use of acetylcysteine for NAI-ALF (and there is no data on its use in remdesivir-associated ALF), both of these cases of remdesivir-associated acute, fulminant ALF showed improvement in ALF with its use in conjunction with discontinuation of potentially hepatotoxic medications without any experiencing any adverse effects.

Conclusion:

The use of the novel antiviral remdesivir in the treatment of COVID-19 pneumonia may put patients at risk of drug-associated acute liver failure. In the two cases presented, the liver failure experienced improved with the administration of acetylcysteine and discontinuation of potentially hepatotoxic medications. More data on the risk of remdesivir-associated acute liver failure and its management with acetylcysteine is needed.

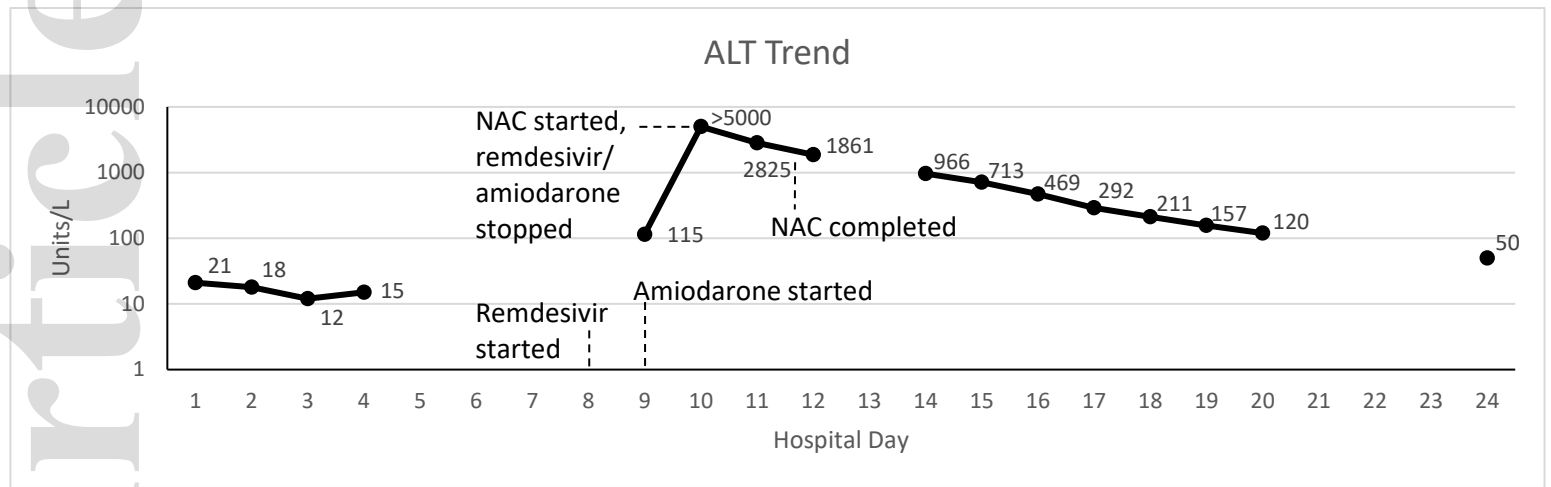
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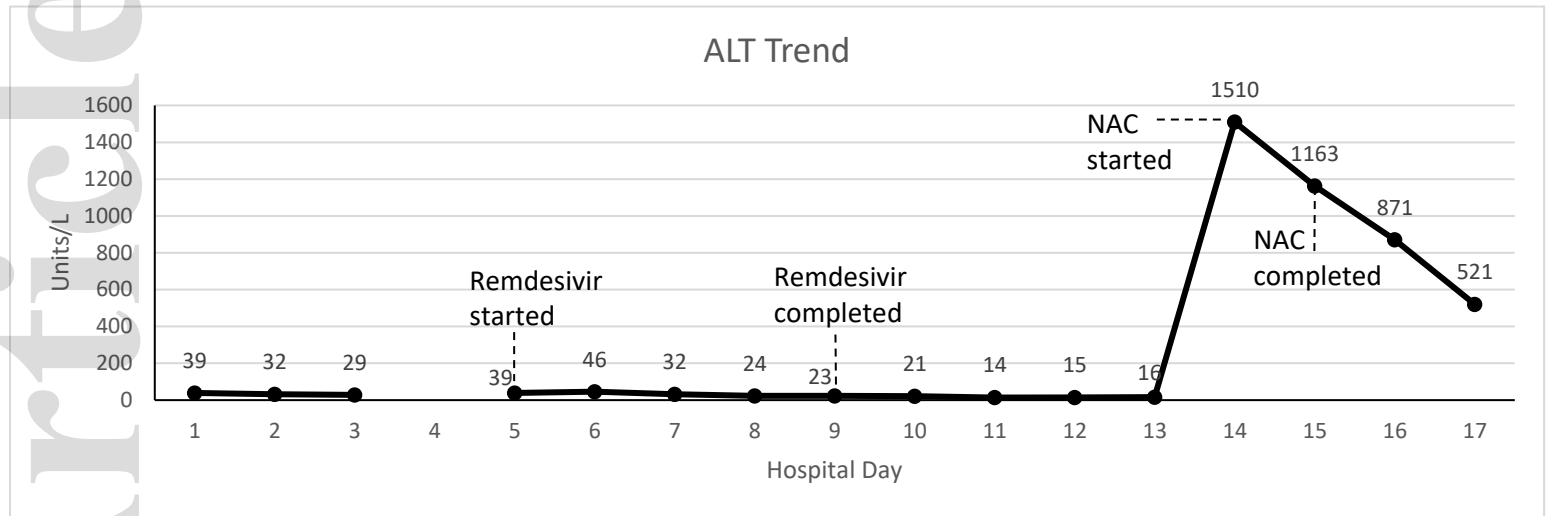
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Figure 1.



Alanine transaminase (ALT) trend by day in case #1 along with significant events (logarithmic trend)

Figure 2.



Alanine transaminase (ALT) trend by day in case #2 along with significant events (linear trend)